



VEDAARA

CLINISAFE

PVT LTD

Pharmacy To Drug Safety

Vedaara Clinisafe is M.P. First Dedicated Institute for Pharmacovigilance & Drug Safety Training.

Our mission is to bridge the gap between Pharmacy education and Industry demand. We provide 6 Weeks intensive, industry-focused training for B.Pharm, D.Pharm, B.Sc and Life Science graduates who want to build careers in PV & Clinical Research.

Global clinical trials industry size is over US\$50 billion. Clinical research and allied sub sectors are rapidly growing industry globally with growth rate of over 12% per annum. As a result of this growth, sector requires highly specialized and skilled professional workforce in the areas of Clinical Research, Clinical Data Management and Pharmacovigilance. Over 10,000 fresher positions are created every year in the clinical trials and allied sectors in India alone. India's share in the clinical trials and related services outsourcing is approx. US\$3 billion. The sector employs over 50,000+ highly skilled professionals. Key global markets like China, South East Asia, Europe and America continue to grow in double digit.

Clinical Research Modules

- ◆ Drug Discovery
- ◆ Overview of Clinical Research
- ◆ Regulations & Guidelines in Clinical Research
- ◆ Roles & Responsibilities of Key Stakeholders
- ◆ Preparation & Planning of Clinical Trials
- ◆ Essential Documentation in Clinical Research & Regulatory Submission
- ◆ Study Start Up Process
- ◆ Clinical Monitoring Essentials
- ◆ Compliance, Auditing & Quality Control in Clinical Research

Pharmacovigilance Modules

- ◆ Overview of Clinical Research
- ◆ Pharmacology – General Principles
- ◆ Introduction to Pharmacovigilance
- ◆ Pharmacovigilance Regulations
- ◆ Hands on Training on Case Processing and Report Generation
- ◆ Safety Reporting and Processing ICSR
- ◆ Aggregate Reports, Signal Detection & Risk Management
- ◆ Documents in Pharmacovigilance
- ◆ Audits & Inspections
- ◆ Advanced Pharmacovigilance & Analytics
- ◆ Artificial Intelligence (AI) in Pharmacovigilance

Clinical Data Management Modules

- ◆ Overview of Clinical Data Management
- ◆ Clinical Data Integrity
- ◆ Data Management Plan
- ◆ Design of Case Report Form
- ◆ CRF Tracking

Entry Level Career Options

- ◆ Clinical Research Coordinator
- ◆ Clinical Trial Assistant
- ◆ Clinical Research Associate
- ◆ Pharmacovigilance Associate
- ◆ Drug Safety Associate
- ◆ Safety Science Analyst
- ◆ Pharmacovigilance Officer
- ◆ Clinical Data Associate
- ◆ Clinical Data Analyst
- ◆ Junior Data Analyst
- ◆ Trainee Junior Data Analyst
- ◆ eTMF Specialist

Mode: eLearning – Live Sessions

Duration: 6 Weeks (Weekends)

Faculty: Experts from the Industry

Eligibility: B.Pharmacy, M.Pharmacy, Pharm.D, Nursing, BSC, MSC, B.Tech Biotech, M.Tech Biotech, BDS, MDS, BHMS, BAMS, BPT, MBBS, Biomedical, Life Sciences

Examination: Online MCQ / Descriptive / Project

Placements: Assured Interviews. Best In the Class Placements in Lifesciences Industry Extensive Placement Support. Pre Placement Training, Pitching Cvs with Recruiters and arranging Interviews with the companies.

Study Material: Recordings, PPT, Notes, Modules, eBooks.

Fee Payment: NEFT / UPI / Card Payment / Cash / Cheque

TOP BENEFITS

• SALARY & GROWTH

- Fresher: 4 LPA - 8 LPA starting
- 2-3 Years: 8 LPA - 15 LPA
- Team Lead: 18 LPA - 25 LPA

• WORK LIFE BALANCE

- WFH Available: TCS, IQVIA, Cognizant
- 9 to 6 Job: No night shift, No Sunday duty
- AC Office: No factory/lab work

• GLOBAL EXPOSURE

- US, EU, India
- Regulations: FDA, EMA, CDSCO
- MNC Culture: TCS, Accenture, Parexel, IQVIA

• LOW STRESS + HIGH DEMAND

• SKILL BASED CAREER

- Software: Argus, ArisG, EudraVigilance
- Knowledge: MedDRA Coding, ICSR, Narrative Writing

TOP RECRUITERS



VEDAARA CLINISAFE PVT. LTD.

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